

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁵ : A61K 31/195</p>	<p>A1</p>	<p>(11) International Publication Number: WO 91/12800 (43) International Publication Date: 5 September 1991 (05.09.91)</p>
<p>(21) International Application Number: PCT/US91/01109 (22) International Filing Date: 27 February 1991 (27.02.91) (30) Priority data: 486,615 28 February 1990 (28.02.90) US PCT/US91/00334 22 January 1991 (22.01.91) WO (34) Countries for which the regional or international application was filed: US et al. (60) Parent Application or Grant (63) Related by Continuation US PCT/US91/00334 (CIP) Filed on 22 January 1991 (22.01.91) (71) Applicant (for all designated States except US): THE UPJOHN COMPANY [US/US]; 301 Henrietta Street, Kalamazoo, MI 49001 (US).</p>		<p>(72) Inventor; and (75) Inventor/Applicant (for US only) : MEGGLASSON, Martin, Durham [-/US]; 5337 Whippoorwill, Kalamazoo, MI 49002 (US). (74) Agent: WRIGHT, Debbie, K.; Corporate Patents & Trademarks, The Upjohn Company, Kalamazoo, MI 49001 (US). (81) Designated States: AT (European patent), AU, BB, BE (European patent), BF (OAPI patent), BG, BJ (OAPI patent), BR, CA, CF (OAPI patent), CG (OAPI patent), CH (European patent), CM (OAPI patent), DE (European patent), DK (European patent), ES (European patent), FI, FR (European patent), GA (OAPI patent), GB (European patent), GR (European patent), HU, IT (European patent), JP, KP, KR, LK, LU (European patent), MC, MG, ML (OAPI patent), MR (OAPI patent), MW, NL (European patent), NO, PL, RO, SD, SE (European patent), SN (OAPI patent), SU, TD (OAPI patent), TG (OAPI patent), US. Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</p>
<p>(54) Title: USE OF 3-GUANIDINOPROPIONIC ACID IN THE TREATMENT AND PREVENTION OF METABOLIC DISORDERS</p> <p>(57) Abstract</p> <p>The present invention provides a method for treating or preventing certain metabolic disorders comprising the administration of 3-guanidinopropionic acid (3-GPA). These metabolic disorders of human and animal metabolism, e.g. hyperglycemia, impaired glucose tolerance, hyperinsulinemia and insulin insensitivity, hyperamylinemia, excess adiposity, and hyperlipidemia, may occur in the following disease states: non-insulin dependent diabetes mellitus (NIDDM), obesity, hypertension and atherosclerosis. 3-GPA may be administered admixed in the diet of farm animals or as a pharmaceutical preparation such as an oral tablet or capsule, by injection, or by implantable sustained release devices thereby increasing the protein content of the carcass while decreasing its fat content. This would produce muscle tissue with less fat.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	ES	Spain	MG	Madagascar
AU	Australia	FI	Finland	ML	Mali
BB	Barbados	FR	France	MN	Mongolia
BE	Belgium	GA	Gabon	MR	Mauritania
BF	Burkina Faso	GB	United Kingdom	MW	Malawi
BG	Bulgaria	GN	Guinea	NL	Netherlands
BJ	Benin	GR	Greece	NO	Norway
BR	Brazil	HU	Hungary	PL	Poland
CA	Canada	IT	Italy	RO	Romania
CF	Central African Republic	JP	Japan	SD	Sudan
CG	Congo	KP	Democratic People's Republic of Korea	SE	Sweden
CH	Switzerland	KR	Republic of Korea	SN	Senegal
CI	Côte d'Ivoire	LI	Liechtenstein	SU	Soviet Union
CM	Cameroon	LK	Sri Lanka	TD	Chad
CS	Czechoslovakia	LU	Luxembourg	TC	Togo
DE	Germany	MC	Monaco	US	United States of America
DK	Denmark				

-1-

USE OF 3-GUANIDINOPROPIONIC ACID IN THE TREATMENT
AND PREVENTION OF METABOLIC DISORDERS

FIELD OF INVENTION

The present invention provides a new use for known compounds. More particularly, the present invention provides a method of treating or preventing certain metabolic disorders of human and animal metabolism, e.g., hyperglycemia, impaired glucose tolerance, hyperinsulinemia, insulin insensitivity, hyperamylinemia, excess adiposity or hyperlipidemia by the administration of 3-guanidinopropionic acid (3-GPA).

BACKGROUND OF THE INVENTION

10 There are several metabolic disorders of human and animal metabolism, e.g., hyperglycemia, impaired glucose tolerance, hyperinsulinemia and insulin insensitivity, hyperamylinemia, excess adiposity, and hyperlipidemia. Some or all of the above disorders may occur in the following disease states: non-insulin dependent diabetes mellitus (NIDDM), obesity, hypertension and atherosclerosis.

15 Hyperglycemia is a condition where the blood glucose level is above the normal level in the fasting state, following ingestion of a meal, or during a provocative diagnostic procedure, e.g., a glucose tolerance test. It can occur in NIDDM as well as obesity. Hyperglycemia can occur without a diagnosis of NIDDM. This condition is called impaired glucose tolerance or pre-diabetes. Impaired glucose tolerance occurs
20 when the rate of metabolic clearance of glucose from the blood is less than that commonly occurring in the general population after a standard dose of glucose has been orally or parenterally administered. It can occur in NIDDM as well as obesity, pre-diabetes and gestational diabetes.

Hyperinsulinemia is defined as having a blood insulin level that is above normal
25 level in the fasting state, following ingestion of a meal or during a provocative diagnostic procedure. It can be seen in NIDDM or obesity and can be associated with or causal in hypertension or atherosclerosis. Hyperinsulinemia can occur without a diagnosis of diabetes. It may occur prior to the onset of NIDDM. Insulin insensitivity, also called insulin resistance, occurs when the insulin-dependent glucose clearance rate is less than
30 that commonly occurring in the general population during diagnostic procedures such as a hyperinsulinemic clamp [See, e.g., DeFronzo, R. A. et al., Am. J. Physiol. 232:E214-E233, (1979)] or a minimal model test. See, e.g., Bergman, R. N. et al., J. Clin. Invest. 68:1456-1467 (1981). Insulin insensitivity is considered also to occur when the

-2-

blood glucose concentration is higher than that commonly occurring in the general population after intravenous administration of insulin (insulin tolerance test) or when the ratio of serum insulin-to-glucose concentration is higher than that commonly occurring in the general population after a 10-16 hour fast. Insulin insensitivity may be found in
5 NIDDM or obesity and can also be associated with or causal to hypertension or atherosclerosis.

Hyperamylinemia is defined as having an abnormally high blood amylin level. Amylin is also known as diabetes associated peptide (DAP) and insulinoma associated polypeptide (IAP). Hyperamylinemia can be seen in NIDDM or obesity.

10 Excess adiposity can be seen in NIDDM associated with obesity as well as obesity without NIDDM. It is defined as a higher fat body mass-to-lean body mass ratio than that commonly occurring in the general population as measured by whole body specific gravity or other generally accepted means.

Hyperlipidemia is defined as having an abnormal level of lipids in the blood.
15 Hyperlipidemia exists when the serum concentration of total cholesterol or total triglycerides or the serum concentration of LDL-cholesterol/HDL-cholesterol is higher than that commonly occurring in the general population. It can be seen in NIDDM or atherosclerosis.

The above disease states could be treated by either ameliorating or preventing the
20 metabolic and biochemical disorders. In addition, humans and animals, which have not been diagnosed as having one of the above disease states but evidencing some or all of the disorders described above, could be benefitted by preventing the development of a currently recognized disease state. Therefore, a compound that is useful in the treatment of hyperglycemia, impaired glucose tolerance, hyperinsulinemia, insulin insensitivity,
25 hyperamylinemia, excess adiposity or hyperlipidemia could also be used to treat or prevent NIDDM, obesity, hypertension or atherosclerosis.

3-Guanidinopropionic acid (3-GPA) is an endogenous metabolite found in animals and humans. See, e.g., Hiraga, Y. et al., J. Chromatography 342:269-275 (1985) and Watanabe, Y. et al., Guanidines, edited by Mori et al., Plenum, New York, pp. 49-58
30 (1983). The compound, which is available from Sigma Chemical Co., has been used extensively in the study of creatine metabolism [See, e.g., Walker, J. B., Adv. Enzymol. 50:177-242 (1979)] and gamma-aminobutyric acid receptor function. See, e.g., Bowery, R. et al., Br. J. Pharmacol. 50:205-218 (1974). Except as noted below, these studies

-3-

do not relate to 3-GPA's utility in treating human or animal disease.

Guanidine, monoguanidine and diguanidine compounds have been shown to produce hypoglycemia. See, e.g., Watanabe, C., J. Biol. Chem. 33:253-265 (1918); Bischoff, F. et al., Guanidine structure and hypoglycemia 81:325-349 (1929). However, these compounds were observed to be toxic. In 1957, biguanide derivatives, e.g. phenformin and metformin, were used clinically as anti-diabetic agents. Some members of this class continue to be used today while others have been withdrawn from the market or banned in the United States and most Western countries. See, e.g., Schafer, G., Diabete Metabol. (Paris) 9:148-163 (1983).

Gamma-guanidinobutyramide also known as Tyformin, and the HCl salt of Tyformin, known as Augmentin, were investigated as potential anti-diabetic agents from the mid-1960's until the mid-1970's. While Augmentin produced hypoglycemia, it was reported to produce hypertension in dogs [See, e.g., Malaisse, W. et al., Horm. Metab. Res. 1:258-265 (1969)] and respiratory and circulatory collapse in rats and rabbits. See, e.g., Buckle, A. et al., Horm. Metab. Res. 3:76-81 (1971). The free acid of the amide was said to lack hypoglycemic activity [See, e.g., Beeson, M. et al., Horm. Metab. Res. 3:188-192 (1971)].

British patent 1,153,424 discloses the use of certain esters and amides of guanidino-aliphatic acids in the treatment of diabetes mellitus where hyperuremia is present. The patent does not disclose that these compounds have an effect on hyperglycemia or any other symptom or pathological state related to diabetes. In a Canadian patent, 891509, the use of esters and amides of guanidinoaliphatic acids were disclosed for treating hyperuremia and hyperglycemia in diabetes mellitus. As noted above, the biologic activity of a guanidino alkanolic acid was known to be different and less favorable so as to be ineffective compared to its amide for treating hyperglycemia.

British patent, 1,195,199 discloses the use of guanidino alkanolic acids or their amides or esters in an insulin-containing, parenterally-administered composition for the treatment of hyperglycemia occurring in diabetes. According to this patent, the combining of a guanidino alkanolic acid, amide or ester with insulin reduces the risk of hypoglycemia as compared to insulin alone. British patent 1,195,200 discloses the use of guanidino alkanolic acids in a composition containing a guanidino alkanolic acid amide or ester derivative for the treatment of hyperglycemia occurring in diabetes. In a subsequent British patent, 1,552,179, the use of guanidino alkanolic acids, their salts,

amides or esters in combination with a gluconeogenesis inhibitor for treating hyperglycemic conditions was disclosed. Metformin was cited as an inhibitor of gluconeogenesis. Biological data indicated that HL 523, the preferred guanidino alkanolic acid derivative, was inactive as a single agent in six of seven experiments where blood glucose concentration was measured in alloxan diabetic mice and only weakly active in the seventh study. Most notably, British patents 1,195,199, 1,195,200 and 1,552,179 do not claim utility for guanidino alkanolic acids, as the sole active component, in compositions for treating hyperglycemic symptoms in diabetes. Among the guanidino alkanolic acids tested, several were inactive as a single agent. Thus, a variety of guanidino alkanolic acids lack significant anti-diabetic activity and combination of these compounds with an agent of known anti-diabetic activity, e.g., metformin, is necessary to show beneficial activity.

Aynsley-Green and Alberti injected rats intravenously with 3-GPA, arginine, guanidine, 4-guanidinobutyramide, and 4-guanidinobutyric acid. Arginine and 3-GPA stimulated insulin secretion transiently, but did not affect the blood glucose concentration while the other compounds stimulated insulin secretion but produced a rise in blood glucose concentration. See, e.g., Aynsley-Green, A. et al., *Horm. Metab. Res.* 6:115-120 (1974). Blachier, et al., observed that 10 mM 3-GPA stimulated insulin secretion by isolated rat islets in vitro. See, e.g., Blachier, F. et al., *Endocrinology* 124:134-141 (1989). The insulin response induced by 3-GPA was 55% of that occurring when arginine was tested at the same concentration. In rats fed a diet supplemented with 10 mg/g 3-GPA for 30-60 days, the heart glycogen content was increased. See, e.g., Roberts, J. et al., *Am. J. Physiol.* 243:H911-H916 (1982). Similarly, skeletal muscle glycogen content was increased in rats fed chow supplemented with 10mg/g of 3-GPA for 6-10 weeks. Mice fed a diet supplemented with 3-GPA at 20 mg/g and supplied with drinking water containing 5 mg/ml 3-GPA for 7-12 weeks had serum glucose concentrations that did not differ significantly from mice receiving unsupplemented chow and water. See, e.g., Moerland, T. et al., *Am. J. Physiol.* 257:C810-C816 (1989).

With respect to adiposity, it is known that in some, but not all cases [See, e.g., Shoubbridge, E. et al., *Biochem. J.* 232:125-131 (1985)], supplementation of the diet with 10-20 mg/g 3-GPA results in decreased body weight. See, e.g., Moerland, *supra* and Mahanna, D. et al., *Exper. Neurol.* 68:114-121 (1980). This effect has been attributed to decreased skeletal muscle mass and has not been attributed to reduced adiposity or

-5-

decreased lipid storage. See, e.g., Mahanna, supra and Shields, R. et al., Lab. Invest. 33:151-158 (1975).

What is needed in the art is a sole therapy to treat or prevent the underlying metabolic disorders in these conditions.

5

INFORMATION DISCLOSURE STATEMENT

The following patents disclosed the use of guanidino-aliphatic acids or their amides or esters for the treatment of insulin-dependent diabetes: British patent 1,153,424; Canadian patent 891509; British patent 1,195,199; British patent 1,195,200; British patent 1,552,179. None of these patents disclosed the use of 3-GPA as a sole therapy, nor as a sole active agent in a pharmaceutical composition. 3-GPA has been shown to stimulate insulin secretion without lowering blood glucose, See, e.g., Ansley-Green, A. et al., Horm. Metab. Res. 6:115-120 (1974) and Blachier, F., Endocrinology 124:134-141 (1989); and to increase heart glycogen content. See, e.g., Roberts, J., Am. J. Physiol. 243:H911-H916 (1982) and Moerland, T., Am. J. Physiol. 257:C810-C816 (1989). It is also known that supplementation of the diet with 3-GPA results in decreased body weight. See Shoubridge, E. A. et al., Biochem. J. 232:125-131 (1985); Moerland, supra; Mahanna, D. A. et al., Exper. Neurol. 68:114-121 (1980); and Shields, R. P. et al., Lab. Invest. 33:151-158 (1975). All of the references cited in this section are discussed above.

20

SUMMARY OF THE INVENTION

The present invention provides a method of treating or preventing a metabolic disorder selected from the group consisting of hyperglycemia, impaired glucose tolerance, hyperinsulinemia, hyperamylinemia, excess adiposity and/or hyperlipidemia in a patient susceptible to or experiencing said disorder comprising the systemic administration of 3-guanidinopropionic acid or a pharmacologically acceptable salt thereof.

Despite extensive suggestions in the literature that it would be ineffective as a sole therapy for insulin-dependent diabetes, surprisingly and unexpectedly, it has been found that 3-GPA causes several biologic effects that are beneficial in the treatment of human disease. It improves plasma glucose level, insulin sensitivity, plasma amylin level, adiposity and plasma lipid level. All of these effects are beneficial in treating NIDDM. 3-GPA offers therapeutic advantage over metformin, a compound currently used to treat NIDDM. See, e.g., Vigneri, R. et al., Diabetes Care 10:118-122 (1987). When

30

-6-

administered to KKA^y (Upjohn) diabetic mice, 3-GPA is more potent and produces a greater decrease in plasma glucose concentration.

3-GPA is ineffective when administered to rodents in a state similar to insulin-dependent diabetes mellitus. 3-GPA did not alter the non-fasting plasma glucose level in lean, normoglycemic C57BL6Job/? mice (Jackson Laboratory) when administered as a 2 mg/g admixture in chow for 13 days. At a higher level, 10 mg/g, 3-GPA did not affect plasma glucose concentration when administered to C57BL6Job/? mice for 4 days, but after administration for 13 days, plasma glucose concentration was decreased. The dose of 3-GPA that produced lower plasma glucose levels in C57BL6Job/? mice was > 6g/kg body weight/day. By contrast, a much lesser dose (130 mg/kg/d) produced an anti-hyperglycemic effect in diabetic KKA^y mice. Since, 3-GPA decreases plasma glucose levels that are elevated, but has little effect at normal plasma glucose concentration, 3-GPA has a great therapeutic benefit in that it ameliorates hyperglycemia with little risk of hypoglycemic reactions in case of overdose.

NIDDM is characterized by hyperglycemia in the fasting or post-prandial state and impaired glucose tolerance after oral or parenteral administration of a glucose solution. 3-GPA, that has been administered to KKA^y mice, a rodent model of NIDDM, decreases the non-fasting plasma glucose concentration and improves glucose tolerance. The minimum effective dose in KKA^y mice is 130 mg/kg/d when administered as an admixture in rodent chow. Higher doses produce a proportionately greater effect. Doses that are less than the minimum effective dose in KKA^y mice may be effective at decreasing blood glucose levels in other species, e.g., human, since elimination is rapid in rodents and may occur more slowly in other species.

Impaired tissue insulin sensitivity and hyperinsulinemia occur in NIDDM [See, e.g., DeFronzo, R., Diabetes 37:667-687 (1988) and Reaven, G., Diabetes 37:1595-607 (1988)], hypertension (See, e.g., Reaven, *supra*), obesity (See, e.g., Glass A., *supra*), and atherosclerosis [See, e.g., Reaven, *supra* and Stout, R. W., Diabetologia 16:141-150 (1979)] and may be etiological factors in these diseases. 3-GPA ameliorates hyperinsulinemia in KKA^y mice and decreases the plasma ratio of insulin-to-glucose concentration, indicating increased insulin sensitivity. Therefore, 3-GPA is useful in the treatment or in the prevention of NIDDM, hypertension, obesity, and atherosclerosis.

Hyperamylinemia may occur in NIDDM, decreasing tissue glucose metabolism [See, e.g., Leighton, B. et al., Nature 335:632-635 (1988)] and altering pancreatic

-7-

hormone secretion [See, e.g., Clark, A., Diabetic Medicine 6:561-567 (1989)]. 3-GPA ameliorates hyperamylinemia and therefore is beneficial in treating disease states in which plasma amylin concentration is increased.

Excess adiposity is an etiological factor in NIDDM and when extreme, represents a disease state in itself. 3-GPA decreases adiposity by decreasing the level of lipids stored in fat and liver tissue. The compound is therefore beneficial in the treatment of obesity alone or in concert with NIDDM. The effect of 3-GPA is selective for lipid-rich tissues (e.g., epididymal fat and fatty liver of ob/ob mice) while muscle mass is unaffected or only minimally affected.

Increased serum low density lipoprotein (LDL) cholesterol concentration is an etiological factor in coronary artery disease. 3-GPA decreases LDL-cholesterol levels in spontaneously hyperlipidemic mice and therefore is useful in treating or preventing hyperlipoproteinemia, atherosclerosis and coronary artery disease.

By sole active pharmaceutical agent is meant that the 3-GPA compound or its salt, administered as claimed herein, is the only pharmaceutical agent in the composition.

By patients susceptible to or experiencing a metabolic disorder, i.e., hyperglycemia, impaired glucose tolerance, hyperinsulinemia, insulin insensitivity, hyperamylinemia, excess adiposity and/or hyperlipidemia is meant a human or animal who exhibits said metabolic disorder and is therefore likely to exhibit one of more of the disease states described above. Such patients are readily diagnosed by a physician or veterinarian of ordinary skill. By treatment is meant the amelioration or total avoidance of the metabolic disorder as described herein. By prevention is meant the avoidance of a currently recognized disease state, as described herein, in a patient evidencing some or all of the metabolic disorders described above.

For all of these purposes, any convenient route of systemic administration is employed, e.g., orally, parenterally, intranasally or intrarectally. In general, the preferred form of administration is orally.

The above compositions may be administered in a sustained release formulation. By sustained release is meant a formulation in which the drug becomes biologically available to the patient at a measured rate over a prolonged period. Such compositions are well-known in the art.

Since 3-GPA decreases body fat without affecting the lean mass, 3-GPA would be of great commercial benefit to the meat, poultry, and fish producing industries in

-8-

achieving its goal of producing leaner animal products. 3-GPA may be administered admixed in the diet of farm animals or as a pharmaceutical preparation such as an oral tablet or capsule, by injection, or by implantable sustained release devices thereby increasing the protein content of the carcass while decreasing its fat content. This would produce muscle tissue with less fat. This benefit of 3-GPA would also impact on the potential health to the meat, poultry, and fish consuming public. The term "farm animals" is defined as animals which are raised for food production. The term includes, but is not limited to, such animals as cattle, poultry, fish, swine, and lamb.

3-GPA increases exercise tolerance in normal mice. Thus the present invention may be useful in treating muscular dysfunction, such as post-poliomyelitis chronic muscle fatigue syndrome or muscular dystrophy, or in treating chronic muscular weakness associated with advanced age or chronic immobilization, or in increasing endurance and exercise in normal humans.

3-GPA also improved the survival rate of mice maintained in a low oxygen environment and therefore is beneficial in treating or preventing disease states involving tissue hypoxia, e.g., peripheral claudication and exercise intolerance in diabetic humans, and angina, myocardial infarction and stroke in diabetic and normal humans.

It is known that glucose-dependent protein crosslinking alters the tertiary structure of several proteins. This protein glycosylation may contribute to diabetic complication and complications of aging in non-diabetic humans, such as neuropathy, nephropathy, retinopathy, hypertension, and atherosclerosis. 3-GPA is useful to block protein glycosylation and therefore be of benefit in treating or preventing this reaction.

The dosage regimen for 3-GPA in accord with this invention will depend on body weight. 3-GPA, in pharmaceutical dosage form, can range from 1-500 mg/kg/day. The preferred dose is 5-100 mg/kg/day. Any sustained released formulations can be used.

DETAILED DESCRIPTION OF THE INVENTION

The present invention is seen more fully by the examples given below.

Example 1 Improvement of Plasma Glucose Level

To test the effect of 3-GPA on non-fasting plasma glucose concentration in KKA^y mice, the mice are administered 3-GPA admixed in rodent chow for 4 days. Plasma is prepared from blood collected by puncturing the retro-orbital sinus of conscious mice that are in the ad libitum fed state. Blood samples are obtained on day 0 (prior to treatment) and on day 4 of treatment. Plasma glucose is measured by an enzyme

-9-

coupled assay using hexokinase and glucose-6-phosphate dehydrogenase. Data is shown as a means \pm S.E.M. for 5 mice per group. Doses that produce a reduction in plasma glucose of 20% or greater are considered to have had a biologically meaningful effect. The percentage reduction in plasma glucose levels by 3-GPA is as follows: 0.8mg/g, 20%; 1.5mg/g, 48%; 2mg/g, 52%; 10mg/g, 77%. The results are summarized in Table I. 3-GPA also decreased the non-fasting plasma glucose level in obese, hyperglycemic C57BL6Job/ob mice which are considered to be a model of NIDDM.

To test the effect of 3-GPA on glucose tolerance in KKA^y mice, the mice are administered 3-GPA admixed in rodent chow at 10 mg/g for 14 days or received unsupplemented chow. Blood samples are collected and plasma analyzed for glucose as described in legend to Table 1. Blood is collected at time 0 (prior to glucose administration) and at intervals after injection of 1.5 g/kg glucose intraperitoneally. Data, as seen in Table II, is shown as means \pm S.E.M. for 5-6 mice per group.

Example 2 Improvement in Insulin Sensitivity

Data supporting the utility of 3-GPA in improving insulin sensitivity and ameliorating hyperinsulinemia is shown in Table III. KKA^y mice are fed chow supplemented with 10mg/g 3-GPA for 14 days or unsupplemented chow. Blood samples are obtained and analyzed for glucose as described in legend to Table I. Plasma insulin is determined using a single antibody radioimmunoassay technique. Data is shown as means \pm S.E.M. for 5-6 mice/group.

Example 3 Improvement of Plasma Amylin Level

KKA^y mice receive 3-GPA as a 2mg/g admixture in chow or unsupplemented chow for 4 days. Ob/ob mice receive 3-GPA as a 2 or 10 mg/g admixture in chow or unsupplemented chow for 30 days. Blood samples are obtained as described in Table I. Amylin is measured in plasma using a double antibody radioimmunoassay. Results are shown in Table IV.

Example 4 Improvement in Adiposity

The effect of 3-GPA on body and organ weights is tested in ob/ob mice. The mice receive 3-GPA as a 2 mg/g or 10 mg/g admixture in chow or are fed unsupplemented chow. Data, which is seen in Table V, is shown as means \pm S.E.M.

Example 5 Improvement in Plasma Lipid Level

Ob/ob mice receive 3-GPA as a 10 mg/g admixture in chow for 13 days or are fed unsupplemented chow. Plasma for lipoprotein analysis is obtained as described in

-10-

Table I. Lipoprotein levels are determined with a Demand Autoanalyzer. Data, which is shown in Table VI, is seen as means \pm S.E.M. for 5 mice per group. For comparison, the plasma lipoprotein profile is shown for 4 lean, non-diabetic C57BL6Jdb/? mice that are untreated.

5 Example 6 Reduced Adiposity and Body Weight in Non-diabetic Obese Mice

3-GPA is tested in A^y mice obtained from Jackson Laboratories (Bar Harbor, Maine). Mice are determined to be free of glycosuria in the fed state using KetoDiaStixTM. 3-GPA is administered for 14 days as an admixture in milled mouse chow at 2 and 5 mg/g chow or unsupplemented chow is provided. Lean and fat body
10 mass is determined using the method of Pace and Rathbun. (See Pace, N. and Rathbun, E.N. Studies on Body Composition. III. The Body Water and Chemically Combined Nitrogen Content Relation to Fat Content. J. Biol. Chem 158:658-691 (1945)). Body weights and the wet weights of excised organs are determined gravimetrically using an analytic laboratory balance.

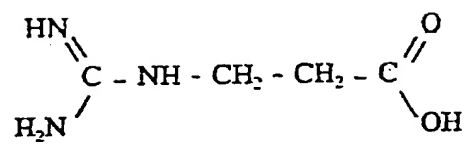
15 As shown in Table 7, 3-GPA significantly decreases the body weight and fat mass of mice as percentages of the body weight, diaphragm and calf muscle, which are selected as representative sources of skeletal muscle, are significantly increased by 3-GPA. The total lean body mass (reflecting primarily skeletal muscle mass) is similarly increased by 3-GPA when expressed as a percentage of the body weight.

20 Example 7 Effect of 3-GPA on Insulin Sensitivity and Body Weight in Adult Rhesus Monkeys

3-GPA is administered orally in gelatin capsule three times daily at a dose of 16 mg/kg for 11 days. Insulin sensitivity is determined using the Bergman Minimal Model technique (Pacini, G. and Bergman, R.N.: MINMOD: a computer program to calculate
25 insulin sensitivity and pancreatic responsivity from the frequently sampled intravenous glucose tolerance test, Computer Meth. Progr. Biomed. 23:113-122, 1986). Insulin sensitivity was improved in all subjects. Body weight decreased in the two heaviest monkeys, but was unaffected in a low body weight monkey. These findings are consistent with previous data from obese rodents where 3-GPA decreased body weight
30 by preferentially lowering the body fat mass without affecting the lean tissue mass and increased insulin sensitivity.

-11-

FORMULA



5

-12-

Table 1

	3-GPA		Non-Fasting Plasma Glucose (mg/dl)	
5	Admixture in Chow (mg/g)	Average dose (g/kg/d)	Days of Treatment	
			0	4
	<u>Experiment 1</u>			
10	Nil	0	481+59	651+52
	2	0.32	441+68	313+75*
15	10	1.74	464+67	148+5**
	*, P=0.006. **,P=0.000.			
20	<u>Experiment 2</u>			
	Nil	0	628+77	536+49
25	0.8	0.13	617+58	429+52*
	1.5	0.21	624+64	279+49**
30	*, P=0.17. **,P=0.006			

-13-

Table 2

		Plasma Glucose (mg/dl)		
5	Time (min)	Control	3-GPA	P-value
	0	484+101	171+21	0.014
10	15	842+83	230+40	0.000
	30	527+81	270+31	0.014
15	60	430+51	190+19	0.002
	120	327+54	153+18	0.013

-14-

Table 3

		Plasma Insulin (microunits/ml)	Plasma Insulin/Glucose (microunits/mg)
5	Control	3,009+419	7.30+1.57
	3-GPA	299+117*	2.04+0.83**
10	*,P=0.000; **,P=0.016		

-15-

Table 4

	Strain	Treatment	Plasma Amylin (ng/ml)
5	KKAY	control	15.0+4.1 (n=5)
		3-GPA	4.3+0.6 (n=3)
10	ob/ob	control	1.3+0.2 (n=6)
		3-GPA (2 mg/g)	1.8+0.2 (n=3)
		3-GPA (10 mg/g)	0.46+0.05 (n=5)
15			

-16-

Table 5

Experiment 1. 31 days treatment. 4-6 mice/group

5	Concentration in Chow:	0	2 mg/g	10 mg/g
	Body Weight (g)	52.3+0.7	49.4+1.1	36.1+1.5
	Liver (g)	4.3+0.2	3.6+0.1	1.6+0.2
10	Epididymal Fat (g)	3.7+0.1	3.8+0.3	2.5+0.2

Experiment 2. 13 days treatment. 5 mice/group

15	Concentration in Chow:	0	10 mg/g
	Body Weight (g)	43.4+1.3	36.8+1.3
20	Liver (g)	2.4+0.2	1.3+0.1
	Heart (g)	0.11+0.01	0.10+0.004
	Diaphragm (g)	0.065+0.004	0.061+0.004
25	Calf muscle (g)	0.050+0.001	0.047+0.003

-17-

Table 6

	Phenotype:	ob/?	ob/ob	ob/ob	P-value
5	3-GPA	(-)	(-)	(+)	ob/ob(-)vs(+)
Cholesterol:					
10	Total	114+4	213+12	214+7	0.932
	Alpha	96+3	194+15	204+9	0.660
15	Beta	18+1	19+4	10+3	0.044
Triglyceride					
20	Total	112+10	170+41	149+25	0.656
	Alpha	70+3	87+4	99+9	0.078
	Beta	42+7	83+38	51+24	0.464
25					

-18-

Table 7

		Addition of 3-GPA to Chow (mg/g):		
		0	2	5
		(n=4)	(n=5)	(n=5)
5	Body Weight (g)			
	Initial	42.72 \pm 1.19	43.55 \pm 1.09	43.02 \pm 0.82
	Final	45.70 \pm 1.64	45.53 \pm 0.97	34.97 \pm 1.16***
10	Fat Mass (g)	13.45 \pm 0.69	13.87 \pm 0.38	8.65 \pm 0.81**
	Lean Body Mass (% BW)	70.60 \pm 0.78	69.54 \pm 0.57	75.47 \pm 1.57*
15	Diaphragm (mg)	97.5 \pm 2.7	103.9 \pm 1.2	96.4 \pm 4.2
	(% BW)	0.21 \pm 0.01	0.23 \pm 0.004	0.28 \pm 0.018**
20	Calf Muscle (mg)	133.5 \pm 23.0	143.2 \pm 5.6	142.5 \pm 4.2
	(%BW)	0.29 \pm 0.05	0.32 \pm 0.018	0.41 \pm 0.018*

25 Data are shown as means \pm S.E.M. Statistical analysis was performed by analysis of variance. P-values for comparison to control mice: *, P<0.05. **, P<0.01. ***, P<0.001.

-19-

Table 8

Subject	Insulin Sensitivity Index		Body Wt. (kg)	
	<u>Placebo</u>	<u>3-GPA</u>	<u>Pre-Tx</u>	<u>3-GPA</u>
5 Monkey 606	0.0040	0.0058	6.6	6.1
Monkey 11	0.0063	0.0076	5.5	5.2
10 Monkey 17	0.0070	0.0110	4.5	4.6

CLAIMS


1. Use of 3-guanidinopropionic acid or a pharmaceutically acceptable salt thereof to prepare a medicament for the treatment of a metabolic disorder consisting of excess
5 adiposity, hyperglycemia, impaired glucose tolerance, hyperinsulinemia, insulin insensitivity, hyperamylinemia, and/or hyperlipidemia.
2. Use of 3-guanidinopropionic acid or a pharmaceutically acceptable salt thereof for the preparation of a medicament for decreasing the fat content and increasing the muscle
10 and protein content of farm animals.
3. A method of treating or preventing a metabolic disorder consisting of excess adiposity, hyperglycemia, impaired glucose tolerance, hyperinsulinemia, insulin insensitivity, hyperamylinemia, and/or hyperlipidemia in a patient susceptible to or
15 experiencing said disorder comprising the systemic administration to said patient of an amount effective to treat or prevent said disorder of 3-guanidinopropionic acid or a pharmaceutically acceptable salt thereof.
4. A method of decreasing the fat content and of increasing the muscle and protein
20 content of farm animals comprising the administration to said farm animal of an amount effective to decrease the fat content and increase the muscle and protein content of 3-guanidinopropionic acid or a pharmaceutically acceptable salt therefor.
5. A method of claim 3 wherein the mode of administration is oral and the patient
25 is a human.
6. A method according to claim 3 wherein said disorder is excess adiposity.
7. A method according to claim 3 wherein said disorder is impaired glucose
30 tolerance.
8. A method according to claim 3 wherein said disorder is hyperinsulinemia.

-21-

9. A method according to claim 3 wherein said disorder is insulin insensitivity.
10. A method according to claim 3 wherein said disorder is hyperglycemia.
- 5 11. A method according to claim 3 wherein said disorder is hyperamylinemia.
12. A method according to claim 3 wherein said disorder is hyperlipidemia.
13. A method according to claim 3 wherein said effective amount of 3-GPA ranges
10 from 1-100 mg/kg/ day.
14. A method according to claim 3 wherein said effective amount of 3-GPA ranges
from 5-100 mg/kg/day.
- 15 16. A method according to claim 4 wherein 3-GPA is administered as an admixture
in the diet, a pharmaceutical preparation, by injection or by implantable sustained
released devices.
17. An oral pharmaceutical composition comprising 3-GPA or a pharmaceutical
20 acceptable salt thereof as the sole active agent and a pharmaceutically acceptable carrier
adapted for humans.
18. A composition of claim 17 wherein the formulation is a sustained release
formulation.

INTERNATIONAL SEARCH REPORT

International Application No PCT/US 91/01109

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) * According to International Patent Classification (IPC) or to both National Classification and IPC IPC ⁵ : A 61 K 31/195						
II. FIELDS SEARCHED Minimum Documentation Searched 7 <table border="1"> <tr> <th>Classification System</th> <th>Classification Symbols</th> </tr> <tr> <td>IPC⁵</td> <td>A 61 K</td> </tr> </table> Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched *			Classification System	Classification Symbols	IPC ⁵	A 61 K
Classification System	Classification Symbols					
IPC ⁵	A 61 K					
III. DOCUMENTS CONSIDERED TO BE RELEVANT *						
Category *	Citation of Document, ** with Indication, where appropriate, of the relevant passages 12	Relevant to Claim No. 13				
X	GB, A, 1195200 (HORLICKS) 17 June 1970 see the whole document, in particular claims 1,2,4,11-21; page 1, lines 22-66 cited in the application ---	1,2,17,18				
X	Metabolism, volume 35, no. 6, June 1986, J.V. Otten et al.: "Thyrotoxic myopathy in mice: accentuation by a creatine transport inhibitor", pages 481-484 see abstract; page 482, tables 1,2 and "results" ---	1,2				
X	The Biochemical Journal, volume 232, no. 1 15 November 1985, The Biochemical Society (London, GB), E.A. Shoubridge et al.: "Biochemical adaptation in the skeletal muscle of rats depleted of creatine with the substrate analogue β -guanidinopropionic acid", pages 125-131 see abstract; page 127, table 1 cited in the application --- ./.	1,2				
* Special categories of cited documents: ** "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "A" document member of the same patent family						
IV. CERTIFICATION						
Date of the Actual Completion of the International Search 17th May 1991		Date of Mailing of this International Search Report 22 JUL 1991				
International Searching Authority EUROPEAN PATENT OFFICE		Signature of Authorized Officer  MISS T. TAZELAAR				

International Application No

PCT/US 91/01109

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category *	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages	Relevant to Claim No.
A	The Journal of Clinical Investigation, volume E61, no. 4, April 1978, D.S. Grosso et al.: "Characterization of a carrier-mediated transport system for taurine in the feta mouse heart in vitro" pages 944-952 see abstract; page 949, table 2 -----	1,2
X	US, A, 3843798 (COOK) 22 October 1974 see claims 1,3; column 4, lines 16-37 -----	17,18

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V. ☒ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE ¹

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. ☒ Claim numbers 3-16 because they relate to subject matter not required to be searched by this Authority, namely:

Please see Rule 39.1 (iv): methods for treatment of the human or animal body by surgery or therapy, as well as diagnostic methods

2. ☐ Claim numbers _____ because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☐ Claim numbers _____ because they are dependent claims and are not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. ☐ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING ²

This International Searching Authority found multiple inventions in this international application as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.
2. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims: _____
3. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers: _____
4. ☐ As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

- ☐ The additional search fees were accompanied by applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

US 9101109

SA 45227

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.
The members are as contained in the European Patent Office EDP file on 12/07/91
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
GB-A- 1195200	17-06-70	None	
US-A- 3843798	22-10-74	None	

EP0 FORM 10079

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82